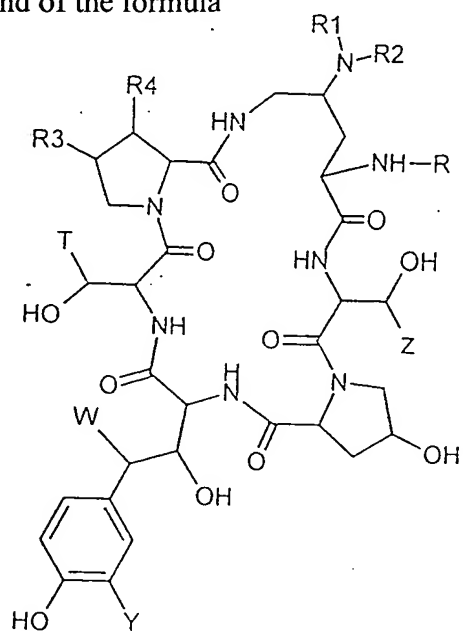


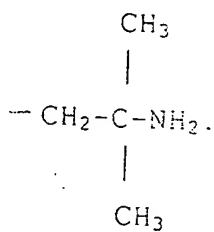
## AMENDMENTS TO THE CLAIMS

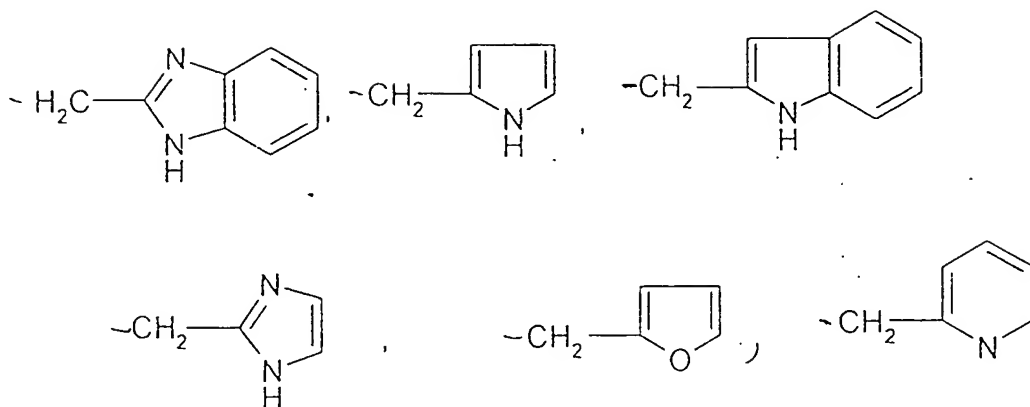
### Claim 1 (currently amended)

A compound selected from the group consisting of all possible isomeric forms and their mixtures, a compound of the formula



either R<sub>1</sub> is hydrogen or methyl and R<sub>2</sub> is selected from the group consisting of cyclohexyl substituted by an amine, -CH<sub>2</sub>-CH<sub>2</sub>NHCH<sub>3</sub>,





$-\text{CHCH}_3\text{CH}_2\text{NH}_2$ ,  $-(\text{CH}_2)_a\text{OH}$  where  $a$  is an integer of 1 to 8,  $-(\text{CH}_2)_b\text{B}-\text{C}\equiv\text{N}$  where

$b$  is an integer of 1 to 8,  $-\text{CHCH}_3\text{C}_6\text{H}_5$ ,  $-(\text{CH}_2)_c-\text{C}(\text{CH}_3)_2\text{NHCOCF}_3$ , and  $-\text{CHCH}_3(\text{CH}_2)_d\text{OH}$  where  $d$  is an integer of 1 to 8

or  $\text{R}_1$  and  $\text{R}_2$  together with the nitrogen to which they are attached form a ring of 3, 4 or 5

carbons optionally substituted by an amine

$\text{R}_3$  is selected from the group consisting of hydrogen, methyl and hydroxyl

$\text{R}_4$  is hydrogen or hydroxyl,

$\text{R}$  is selected from the group consisting of alkyl and cycloalkyl of up to 30 carbon atoms, optionally containing at least one heteroatom, at least one heterocycle and alkyl or cyclic acyl of up to 30 carbon atoms optionally containing at least one heteroatom, and/or at least one heterocycle,

$\text{T}$  is selected from the group consisting of hydrogen, methyl,  $-\text{CH}_2\text{CONH}_2$ ,  $-\text{CH}_2\text{C}\equiv\text{N}$ , and  $-(\text{CH}_2)_2\text{NH}_2$  and  $-(\text{CH}_2)_2\text{N}(\text{alk})^+\text{X}^-$ ,  $\text{X}$  is halogen and alk is alkyl of up to 8 carbon atoms,

$\text{Y}$  is selected from the group consisting of hydrogen, hydroxyl, halogen and  $-\text{OSO}_3\text{H}$  or a salt thereof,

W is hydrogen or OH,

Z is hydrogen or methyl and its non-toxic, pharmaceutically acceptable acid addition salt.

**Claim 2** (currently amended)

A The compound of claim 1 in which T is hydrogen.

**Claim 3** (currently amended)

A The compound of claim 1 in which W is hydrogen.

**Claim 4** (currently amended)

A The compound of claim 1 in which Z is methyl.

**Claim 5** (currently amended)

A The compound of claim 1 in which Y is hydrogen.

**Claim 6** (currently amended)

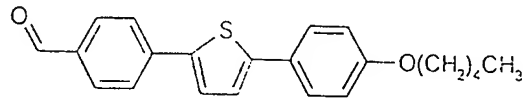
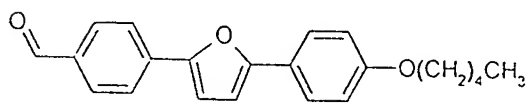
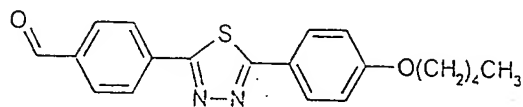
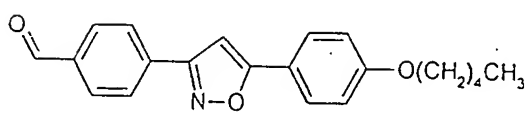
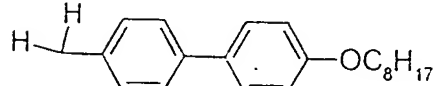
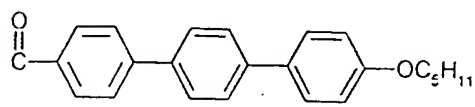
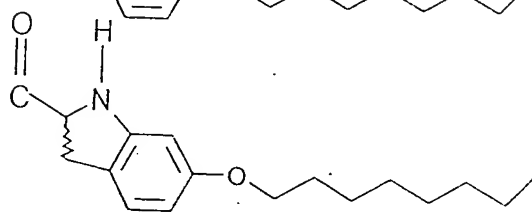
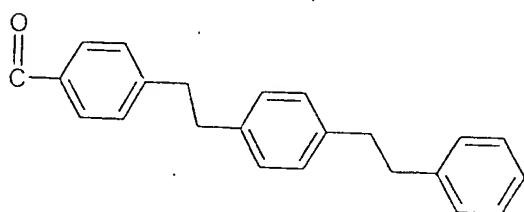
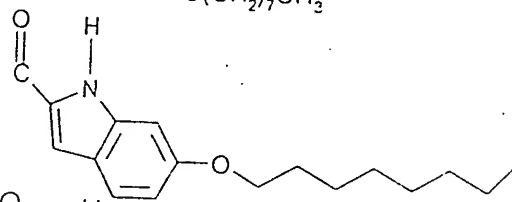
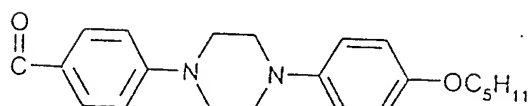
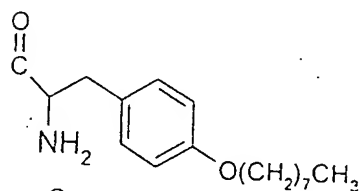
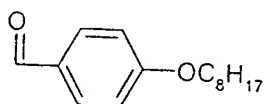
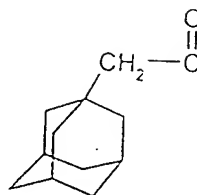
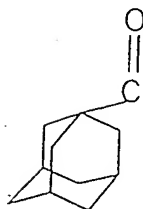
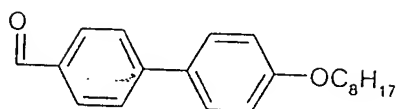
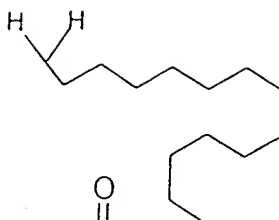
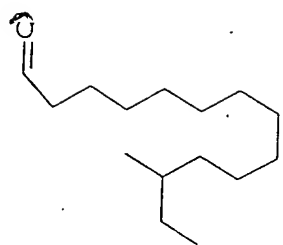
A The compound of claim 1 in which R<sub>3</sub> is methyl.

**Claim 7** (currently amended)

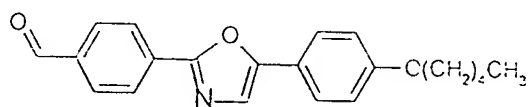
A The compound of claim 1 in which R<sub>4</sub> is hydroxyl.

**Claim 8** (currently amended)

A The compound of claim 1 in which R is selected from the group consisting of

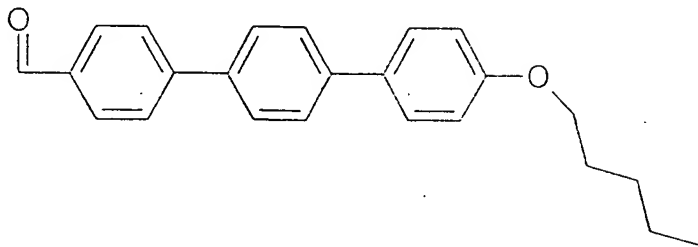


and



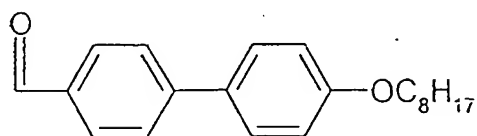
**Claim 9** (currently amended)

A The compound of claim 8 in which R is



**Claim 10** (currently amended)

A The compound of claim 8 in which R is

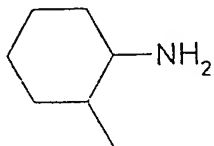


**Claim 11** (currently amended)

A The compound of claim 1 in which R<sub>1</sub> is hydrogen.

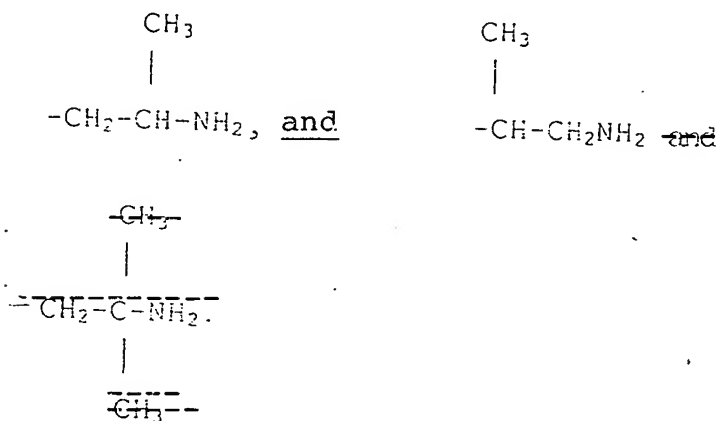
**Claim 12** (currently amended)

A The compound of claim 1 in which R<sub>2</sub> is



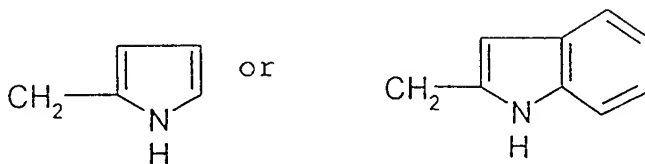
**Claim 13** (currently amended)

A The compound of claim 1 in which R<sub>2</sub> is selected from the group consisting of



**Claim 14** (currently amended)

A The compound of claim 1 in which R<sub>2</sub> is



**Claim 15** (currently amended)

A The compound of claim 1 selected from the group consisting of

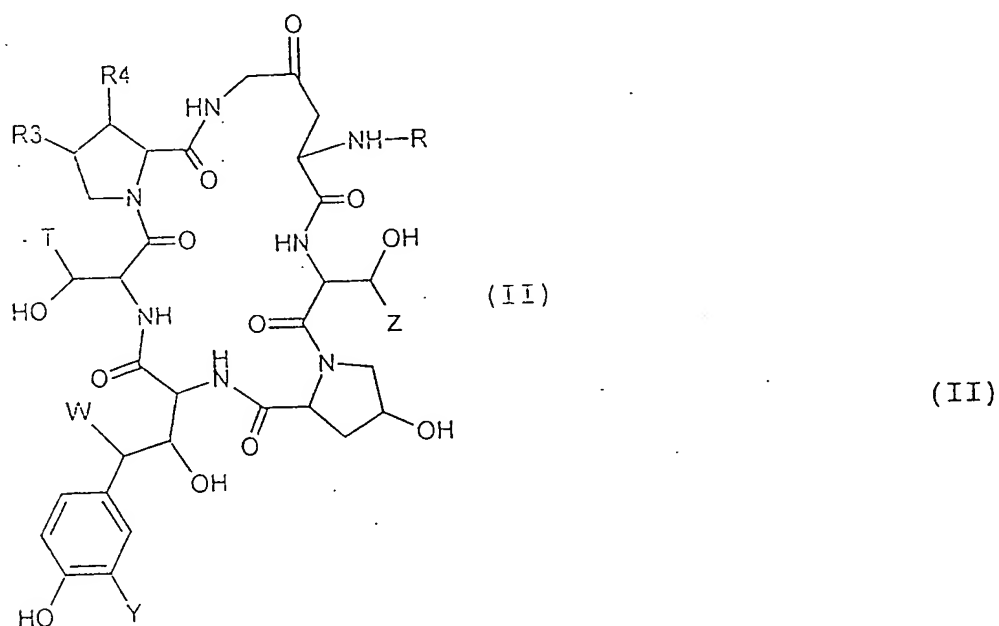
- 1-[4-[[[(1H-benzimidazol-2-yl)-methyl)-amino-N2-[[4''-(pentyloxy) [1,2':4', 1''-

terphenyl]-4-yl]-carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]5-L-serine-  
echinocandine B trifluoroacetate (isomer B), and

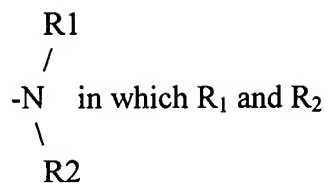
- trans 1-[4-[(2-aminocyclo-hexyl)-amino]-N2-[[4''(pentyloxy) [1,1':4', 1''-terphenyl]-4-yl]-carbonyl]-L-ornithine]-4-[4-hydroxyphenyl)-L-threonine]-5-L-serine-  
echinocandine B trifluoroacetate (isomer A).

**Claim 16 (currently amended)**

A process for the preparation of a compound of claim 1 comprising reacting a  
compound of the formula



wherein R, R<sub>3</sub>, R<sub>4</sub>, T, Y, W and Z are defined as in claim 1 with an amine or amino  
derivative compound capable of introducing



are defined as in claim 1 and optionally to the action of a reducing agent  
and/or an amine functionalization agent,  
and/or an acid to form the salt of the product of claim 1,  
and/or a separation agent of the different isomers obtained.

**Claims 17 and 18** (cancelled)

**Claim 19** (currently amended)

An antifungal composition comprising an antifungally effective amount of a  
compound of claim  $\pm$  15 and an inert pharmaceutical carrier.

**Claim 20** (currently amended)

A method of treating fungal infections in warm-blooded animals comprising  
administering to warm-blooded animals in need thereof an antifungally effective  
amount of a compound of claim  $\pm$  15.